### **AMENDMENT**

## IN THE SPECIFICATION:

Page 1, amend lines 1-5 as follows:

SONY International (Europe) GmbH S30156(M)

## TITLE OF THE INVENTION

Selective Metallisation of Nucleic Acids via Metal Nanoparticles Produced In Situ

## FIELD OF THE INVENTION

Page 1, add line 7 as follows:

#### **BACKGROUND OF THE INVENTION**

Page 3, after line 25, add the following:

# **OBJECTS AND SUMMARY OF THE INVENTION**

Page 7, after line 21, insert the following:

## BRIEF DESCRIPTION OF THE DRAWINGS

The invention will now be described in further detail with respect to the accompanying figures in which:

Figure 1 shows the UV-visible absorption spectra of the Pt(II)-terpyridine-DNA conjugate and the Pt-DNA composites produced according to Example 1.

Figure 2 shows an AFM image of a Pt-DNA composite produced according to Example 1 before treatment with a solution of GoldEnhance® according to Example 4.

Figure 3 shows an AFM image of a Pt-DNA composite produced according to Example 1 after treatment with a solution of GoldEnhance® according to Example 4.

Figure 4 shows an AFM image of another spot of the sample shown in Figure 3.

Figure 5 shows an AFM image of a Pt-DNA composite produced according to Example 2 before treatment with a solution of GoldEnhance® according to Example 5.

Figure 6 shows an AFM image of a Pt-DNA composite produced according to Example 2 after treatment with a solution of GoldEnhance® according to Example 6.

Figure 7 shows the most likely positions for "metalation" at the N-7 atoms of the purine nucleotides (G and A) of a nucleic acid.

Figure 8 shows several variations of metal (M) – ligand  $(L^1, L^2, \text{ and } L^3, X \text{ or } Z)$  complexes, (the charges have been omitted for simplicity).

Figure 9 schematically shows metalation of specific bases within oligonucleotide subunits at sites that are inherently present, (the charges have been omitted for simplicity);

Figure 10 schematically shows metalation of specific bases within oligonucleotide subunits at sites that have been introduced by chemical modification; (the charges have been omitted for simplicity).

Figure 11 shows examples of substitution-inert metal (M) complexes attached to nucleic acid interacting groups of the general formula INT-CON-LIG-M(L)<sub>n</sub>.

Figure 12 schematically shows the covalent attachment of substitution-inert metal complexes to specific bases within oligonucleotide subunits, before or after hybridization at complementary segments of longer nucleic acids; (the charges have been omitted for simplicity).

Figure 13 shows an AFM image of an unmodified non-platinated DNA after treatment with a solution of GoldEnhance®.

Figure 14 shows an AFM image of an unmodified non-platinated DNA after treatment with a solution of GoldEnhance®.

# **DETAILED DESCRIPTION OF THE INVENTION**

Page 13, line 25, to page 15, line 4:

The invention will now be described in further detail with respect to the accompanying figures in which

Figure 1 shows the UV-visible absorption spectra of the Pt(II) terpyridine DNA conjugate and the Pt-DNA composites produced according to Example 1.

Figure 2 shows an AFM image of a Pt-DNA composite produced according to Example 1 before treatment with a solution of GoldEnhance® according to Example 4.

Figure 3 shows an AFM image of a Pt-DNA composite produced according to Example 1 after treatment with a solution of GoldEnhance® according to Example 4.

Figure 4 shows an AFM image of another spot of the sample shown in Figure 3.

Figure 5 shows an AFM image of a Pt-DNA composite produced according to Example 2 before treatment with a solution of GoldEnhance® according to Example 5.

Figure 6 shows an AFM image of a Pt-DNA composite produced according to Example 2 after treatment with a solution of GoldEnhance® according to Example 6.

Figure 7 shows the most likely positions for "metalation" at the N-7 atoms of the purine nucleotides (G and A) of a nucleic acid;

Figure 8 shows several variations of metal (M) ligand (L<sup>1</sup>, L<sup>2</sup>, and L<sup>3</sup>, X or Z) complexes, (the charges have been omitted for simplicity);

Figure 9 schematically shows metalation of specific bases within oligonucleotide subunits at sites that are inherently present, (the charges have been omitted for simplicity);

Figure 10 schematically shows metalation of specific bases within oligonucleotide subunits at sites that have been introduced by chemical modification; (the charges have been omitted for simplicity);

Figure 11 shows examples of substitution inert metal (M) complexes attached to nucleic acid interacting groups of the general formula INT CON LIG M(L)<sub>n</sub>;

Figure 12 schematically shows the covalent attachment of substitution inert metal complexes to specific bases within oligonucleotide subunits, before or after hybridization at complementary segments of longer nucleic acids; (the charges have been omitted for simplicity); and; Figure 13 shows an AFM image of an unmodified non-platinated DNA after treatment with a solution of GoldEnhance®.

Figure 14 shows an AFM image of an unmodified non-platinated DNA after treatment with a solution of GoldEnhance®.